IN THE CLAIMS:

- 1. (currently amended) Use of an antimuscarinic agent for the manufacture of a medicament for oral administration of a pharmaceutically effective dose of the antimuscarinic agent when needed to a mammal with unstable or overactive bladder, A method for treating unstable or overactive urinary bladder in a mammal, said method comprising orally administering to a mammal a pharmaceutically effective dose of an antimuscarinic agent and when needed, whereby a symptomatic relief of urgency and/or frequency is achieved.
- 2. (currently amended) The <u>use method</u> as claimed in claim 1, wherein the antimuscarinic agent is one or more compounds selected from tolterodine and related compounds.
- 3. (currently amended) The $\frac{method}{method}$ according to claim 2, wherein the compound is tolterodine or a pharmaceutically acceptable salt thereof.
- 4. (currently amended) The use method according to claim 1, wherein the antimuscarinic agent is selected from oxybutynin, darifenacin, solifenacin, and the pharmaceutically acceptable salts and derivatives thereof.
- 5. (currently amended) The $\frac{method}{method}$ according to any of claims 1 to 4, wherein the mammal is human.
- 6. (currently amended) The $\frac{method}{method}$ according to claim 5, wherein the pharmaceutically effective dose is 2 mg or 4 mg

of the antimuscarinic agent, administered as a controlled release tablet or capsule.

- 7. (currently amended) The use method according to claim 5, wherein two pharmaceutically effective doses of the antimuscarinic agent are administered daily at an interval of within 8-12 hours.
- 8. (currently amended) The use method according to claim 7, wherein the pharmaceutically effective dose is 1 mg of the antimuscarinic agent, administered as an immediate release tablet or capsule.
- 9. (currently amended) The <u>use method</u> according to claim 7, wherein the pharmaceutically effective dose is 1 mg or 2 mg of the antimuscarinic agent, administered as a controlled release tablet or capsule.
- 10. (currently amended) The $\frac{\text{method}}{\text{method}}$ according to any of claims 1-5 or 7-9, wherein the pharmaceutically effective doses of the antimuscarinic agent are taken within the interval of 8 hours.
- 11. (currently amended) The use method according to any of claims 1-5 or 7-9, wherein the pharmaceutically effective doses of the antimuscarinic agent are taken within the interval of 9 hours.
- 12. (currently amended) The use method according to any of claims 1-5 or 7-9, wherein the pharmaceutically effective

doses of the antimuscarinic agent are taken within the interval of 10 hours.

- 13. (currently amended) The use method according to any of claims 1-5 or 7-9, wherein the pharmaceutically effective doses of the antimuscarinic agent are taken within the interval of 11 hours.
- 14. (currently amended) The use method according to any of claims 1-5 or claims 7-9, wherein the pharmaceutically effective doses of the antimuscarinic agent are taken within the interval of 12 hours.

15. (canceled)